Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

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- 1. (Original) A pharmaceutical granulated product having improved granulatability, which contains a pharmaceutical compound with poor wettability and a surfactant.
- 2. (Original) A granulated product, wherein the product contains a compound with poor wettability and a surfactant, and at least about 35% by weight with respect to the total weight of the product does not pass through a 100-mesh sieve.
- 3. (Original) The granulated product according to claim 2, wherein the weight ratio of the compound and the surfactant is 1: about 0.001 to about 2.
- 4. (Original) The granulated product according to claim 3, wherein the weight ratio is 1: about 0.001 to less than 1.
- 5. (Original) The granulated product according to claim 3, wherein the weight ratio is 1: about 0.001 to less than 0.1.
- 6. (Original) The granulated product according to claim 3, wherein the weight ratio is 1 : about 0.005 to about 0.05.
- 7. (Original) The granulated product according to claim 2, wherein the ratio of the compound with respect to the total granulated product is about 20% by weight or more.
- 8. (Original) The granulated product according to claim 2, wherein the compound is a pharmaceutical compound.
- 9. (Currently amended) A molded product made by molding the granulated product according to any one of claims 2 to 8 claim 2.
- 10. (Original) A method for improving granulatability of a pharmaceutical composition containing a pharmaceutical compound with poor wettability, which comprises adding a surfactant before or during the granulation.
- 11. (Original) A method for preparing a granulated product containing a compound with poor wettability, having improved granulatability, which comprises adding a surfactant

in the weight ratio of about 0.001 to about 2 with respect to the compound before or during the granulation.

- 12. (Original) The method according to claim 11, wherein a granulated product is obtained in which at least about 35% by weight with respect to the total weight of the product does not pass through a 100-mesh sieve.
- 13. (Original) The method according to claim 11, which involves wet granulation in a binder solution containing a surfactant.
- 14. (Original) The method according to claim 13, wherein the concentration of the surfactant in the binder solution is about 1 to about 1,000 mmol/L.
- 15. (Original) The method according to claim 13, wherein the concentration of the surfactant in the binder solution is about 10 to about 100 mmol/L.
- 16. (Original) The method according to claim 11, wherein the compound is a pharmaceutical compound.
- 17. (Currently amended) A method for preparing a molded product, comprising molding the granulated product which is obtained by the method according to any one of claims 11 to 16 claim 11.
 - 18. (Canceled)

19. (Canceled)